US Environmental Protection Agency
Office of Pesticide Programs

Pesticide Registration Improvement Renewal Act – PRIA II
Fee Table – Effective October 1, 2007

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EPA Number of Fee Category (EPA No.), corresponding number in the Congressional Record of July 21, 2007 (CR No., Title of Fee Category (Action), Decision Time Review Periods for FY 08, 09 and 10 in months, and the Registration Service Fee for FY08

EPA No.	CR No.	Action	Decision time (months) FY 08	FY 09	FY 10	FY 08 Registration Service Fee ($)
TABLE 1.		**Registration Division — New Active Ingredients**				
R010	1	Food use **(1)**	24	24	24	516,300
R020	2	Food use; reduced risk **(1)**	18	18	18	516,300
R030	3	Food use; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit and temporary tolerance same as #R040 **(1)**	24	24	24	570,700
R040	4	Food use; Experimental Use Permit application; establish temporary tolerance; submitted before application for registration; credit $326,025 toward new active ingredient application that follows	18	18	18	380,500
R050	5	Food use; application submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit and temporary tolerance are granted **(1)**	14	14	14	190,300
R060	6	Non-food use; outdoor **(1)**	21	21	21	358,700
R070	7	Non-food use; outdoor; reduced risk **(1)**	16	16	16	358,700
R080	8	Non-food use; outdoor; Experimental Use Permit application submitted simultaneously with application for registration; decision time for Experimental Use Permit same as #R090 **(1)**	21	21	21	396,800
R090	9	Non-food use; outdoor; Experimental Use Permit application submitted before application for registration; credit $228,225 toward new active ingredient application that follows	16	16	16	266,300
R100	10	Non-food use; outdoor; submitted after Experimental Use Permit application; decision time begins after Experimental Use Permit is granted **(1)**	12	12	12	130,500
R110	11	Non-food use; indoor **(1)**	20	20	20	199,500
R120	12	Non-food use; indoor; reduced risk **(1)**	14	14	14	199,500
R121	13	Non-food use; indoor; Experimental Use Permit application submitted before application for registration; credit $100,000 toward new active ingredient application that follows	18	18	18	150,000
R122	14	Enriched isomer(s) of registered mixed-isomer active ingredient **(1)**	18	18	18	260,900
R123	15	Seed treatment only; includes non-food and food uses; limited uptake into Raw Agricultural Commodities **(1)**	18	18	18	388,200
R124	16	Conditional Ruling on Preapplication Study Waivers; applicant-initiated	6	6	6	2,080

TABLE 2.		**Registration Division — New Uses**				
R130	17	First food use; indoor; food/food handling **(1)**	21	21	21	157,500
R140	18	Additional food use; Indoor; food/food handling	15	15	15	36,750
R150	19	First food use **(1)**	21	21	21	217,400
R160	20	First food use; reduced risk **(1)**	16	16	16	217,400
R170	21	Additional food use	15	15	15	54,400
R180	22	Additional food use; reduced risk	10	10	10	54,400
R190	23	Additional food uses; 6 or more submitted in one application	15	15	15	326,400
R200	24	Additional food uses; 6 or more submitted in one application; reduced risk	10	10	10	326,400
R210	25	Additional food use; Experimental Use Permit application; establish temporary tolerance; no credit toward new use registration	12	12	12	40,300
R220	26	Additional food use; Experimental Use Permit application; crop destruct basis; no credit toward new use registration	6	6	6	16,320
R230	27	Additional use; non-food; outdoor	15	15	15	21,740
R240	28	Additional use; non-food; outdoor; reduced risk	10	10	10	21,740
R250	29	Additional use; non-food; outdoor; Experimental Use Permit application; no credit toward new use registration	6	6	6	16,320
R260	30	New use; non-food; indoor	12	12	12	10,500
R270	31	New use; non-food; indoor; reduced risk	9	9	9	10,500
R271	32	New use; non-food; indoor; Experimental Use Permit application; no credit toward new use registration	6	6	6	8,000
R272	33	Review of Study Protocol; applicant-initiated; excludes DART, pre-registration conferences, Rapid Response review, DNT protocol review, protocols needing HSRB review	3	3	3	2,080
R273	34	Additional use; seed treatment; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food or non-food uses	12	12	12	41,500
R274	35	Additional uses; seed treatment only; 6 or more submitted in one application; limited uptake into Raw Agricultural Commodities; includes crops with established tolerances (e.g., for soil or foliar application); includes food and/or non-food uses	12	12	12	249,000
TABLE 3.		**Registration Division — Import And Other Tolerances**				
R280	36	Establish import tolerance; new active ingredient or first food use **(1)**	21	21	21	262,500
R290	37	Establish import tolerance; additional food use	15	15	15	52,500
R291	38	Establish import tolerances; additional food uses; 6 or more crops submitted in one petition	15	15	15	315,000

R292	39	Amend an established tolerance (e.g., decrease or increase); domestic or import; applicant-initiated	10	10	10	37,300
R293	40	Establish tolerance(s) for inadvertent residues in one crop; applicant-initiated	12	12	12	44,000
R294	41	Establish tolerances for inadvertent residues; 6 or more crops submitted in one application; applicant-initiated	12	12	12	264,000
R295	42	Establish tolerance(s) for residues in one rotational crop in response to a specific rotational crop application; applicant-initiated	15	15	15	54,400
R296	43	Establish tolerances for residues in rotational crops in response to a specific rotational crop petition; 6 or more crops submitted in one application; applicant-initiated	15	15	15	326,400
TABLE 4.		**Registration Division — New Products**				
R300	44	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	3	1,300
R301	45	New product; identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy, where applicant does not own all required data and does not have a specific authorization letter from data owner.	4	4	4	1,560
R310	46	New end-use or manufacturing-use product; requires review of data package within RD; includes reviews and/or waivers of data for only: • product chemistry and/or • acute toxicity and/or • public health pest efficacy	6	6	6	4,360
R311	49	New product; requires approval of new food-use inert; applicant-initiated; excludes approval of safeners	12	12	12	15,540
R312	50	New product; requires approval of new non-food-use inert; applicant-initiated	6	6	6	8,300
R313	51	New product; requires amendment to existing inert tolerance exemption (e.g., adding post-harvest use); applicant-initiated	10	10	10	11,420

R320	47	New product; new physical form; requires data review in science divisions	12	12	12	10,880
R330	48	New manufacturing-use product; registered active ingredient; selective data citation	12	12	12	16,320
R331	52	New product; repack of identical registered end-use product as a manufacturing-use product; same registered uses only	3	3	3	2,080
R332	53	New manufacturing-use product; registered active ingredient; unregistered source of active ingredient; submission of completely new generic data package; registered uses only	24	24	24	233,000
TABLE 5.		**Registration Division — Amendments To Registration**				
R340	54	Amendment requiring data review within RD (e.g., changes to precautionary label statements, or source changes to an unregistered source of active ingredient) **(2)**	4	4	4	3,280
R350	55	Amendment requiring data review in science divisions (e.g., changes to REI, or PPE, or PHI, or use rate, or number of applications; or add aerial application; or modify GW/SW advisory statement) **(2)**	8	8	8	10,880
R370	56	Cancer reassessment; applicant-initiated	18	18	18	163,100
R371	57	Amendment to Experimental Use Permit; requires data review / risk assessment	6	6	6	8,300
R372	58	Refined ecological and/or endangered species assessment; applicant-initiated	18	18	12	155,300
TABLE 6.		**Antimicrobials Division — New Active Ingredients**				
A380	59	Food use; establish tolerance exemption **(1)**	24	24	24	94,500
A390	60	Food use; establish tolerance **(1)**	24	24	24	157,500
A400	61	Non-food use; outdoor; FIFRA §2(mm) uses **(1)**	18	18	18	78,750
A410	62	Non-food use; outdoor; uses other than FIFRA §2(mm) **(1)**	21	21	21	157,500
A420	63	Non-food use; indoor; FIFRA §2(mm) uses **(1)**	18	18	18	52,500
A430	64	Non-food use; indoor; uses other than FIFRA §2(mm) **(1)**	20	20	20	78,750
A431	65	Non-food use; indoor; low-risk and low-toxicity food-grade active ingredient(s); efficacy testing for public health claims required under GLP and following DIS/TSS or AD-approved study protocol	12	12	12	55,000
TABLE 7.		**Antimicrobials Division — New Uses**				
A440	66	First food use; establish tolerance exemption **(1)**	21	21	21	26,250
A450	67	First food use; establish tolerance **(1)**	21	21	21	78,750
A460	68	Additional food use; establish tolerance exemption	15	15	15	10,500
A470	69	Additional food use; establish tolerance	15	15	15	26,250

A480	70	Additional use; non-food; outdoor; FIFRA §2(mm) uses	9	9	9	15,750
A490	71	Additional use; non-food; outdoor; uses other than FIFRA §2(mm)	15	15	15	26,250
A500	72	Additional use; non-food; indoor; FIFRA §2(mm) uses	9	9	9	10,500
A510	73	Additional use; non-food; indoor; uses other than FIFRA §2(mm)	12	12	12	10,500
A520	74	Experimental Use Permit application	9	9	9	5,250
A521	75	Review of public health efficacy study protocol within AD; per AD Internal Guidance for the Efficacy Protocol Review Process; applicant-initiated; Tier 1	6	4	3	2,000
A522	76	Review of public health efficacy study protocol outside AD by members of AD Efficacy Protocol Review Expert Panel; applicant-initiated; Tier 2	18	15	12	10,000
TABLE 8.		**Antimicrobials Division — New Products & Amendments**				
A530	77	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	3	1,050
A531	78	New product; identical or substantially similar in composition and use to a registered product; registered source of active ingredient; selective data citation only for data on product chemistry and/or acute toxicity and/or public health pest efficacy, where applicant does not own all required data and does not have a specific authorization letter from data owner.	4	4	4	1,500
A532	85	New product; identical or substantially similar in composition and use to a registered product; registered active ingredient; unregistered source of active ingredient; cite-all data citation except for product chemistry; product chemistry data submitted	4	4	4	4,200
A540	79	New end use product; FIFRA §2(mm) uses only	4	4	4	4,200
A550	80	New end-use product; uses other than FIFRA §2(mm); non-FQPA product	6	6	6	4,200
A560	81	New manufacturing-use product; registered active ingredient; selective data citation	12	12	12	15,750
A570	82	Label amendment requiring data submission **(2)**	4	4	4	3,150
A571	83	Cancer reassessment; applicant-initiated	18	18	18	78,750

A572	84	Refined ecological risk and/or endangered species assessment; applicant-initiated	18	18	12	75,000
TABLE 9.		**Biopesticide & Pollution Prevention Division — Microbial & Biochemical Pesticides; New Products & Amendments**				
B580	86	New active ingredient; food use; establish tolerance **(1)**	18	18	18	42,000
B590	87	New active ingredient; food use; establish tolerance exemption **(1)**	16	16	16	26,250
B600	88	New active ingredient; non-food use **(1)**	12	12	12	15,750
B610	89	Food use; Experimental Use Permit application; establish temporary tolerance exemption	9	9	9	10,500
B620	90	Non-food use; Experimental Use Permit application	6	6	6	5,250
B621	91	Extend or amend Experimental Use Permit	6	6	6	4,200
B630	92	First food use; establish tolerance exemption	12	12	12	10,500
B631	93	Amend established tolerance exemption	9	9	9	10,500
B640	94	First food use; establish tolerance **(1)**	18	18	18	15,750
B641	95	Amend established tolerance (e.g., decrease or increase)	12	12	12	10,500
B650	96	New use; non-food	6	6	6	5,250
B660	97	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	3	1,050
B670	98	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product specific data or with request for data waivers supported by scientific rationales	6	6	6	4,200
B671	99	New product; food use; unregistered source of active ingredient; requires amendment of established tolerance or tolerance exemption; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	16	16	16	10,500
B672	100	New product; non-food use or food use having established tolerance or tolerance exemption; unregistered source of active ingredient; no data compensation issues; all Tier I data requirements for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product-specific data or with request for data waivers supported by scientific rationales	12	12	12	7,500

B680	101	Label amendment requiring data submission **(2)**	4	4	4	4,200
B681	102	Label amendment; unregistered source of active ingredient; supporting data require scientific review	6	6	6	5,000
B682	103	Protocol review; applicant-initiated; excludes time for HSRB review (pre application)	3	3	3	2,000
TABLE 10.		**Biopesticide & Pollution Prevention Division — Straight Chain Lepidopteran Pheromones (Sclps)**				
B690	104	New active ingredient; food or non-food use **(1)**	6	6	6	2,100
B700	105	Experimental Use Permit application; new active ingredient or new use	6	6	6	1,050
B701	106	Extend or amend Experimental Use Permit	3	3	3	1,050
B710	107	New product; identical or substantially similar in composition and use to a registered product; no data review or only product chemistry data; cite-all data citation, or selective data citation where applicant owns all required data, or applicant submits specific authorization letter from data owner. Category also includes 100% re-package of registered end-use or manufacturing-use product that requires no data submission nor data matrix.	3	3	3	1,050
B720	108	New product; registered source of active ingredient; all Tier I data for product chemistry, toxicology, non-target organisms, and product performance must be addressed with product specific data or with request for data waivers supported by scientific rationales	4	4	4	1,050
B721	109	New product; unregistered source of active ingredient	6	6	6	2,200
B722	110	New use and/or amendment to tolerance or tolerance exemption	6	6	6	2,200
B730	111	Label amendment requiring data submission **(2)**	4	4	4	1,050
TABLE 11.		**Biopesticide & Pollution Prevention Division — Plant Incorporated Protectants (Pips)**				
B740	112	Experimental Use Permit application; registered active ingredient; non-food/feed or crop destruct basis; no SAP review required **(3)**	6	6	6	78,750
B750	113	Experimental Use Permit application; registered active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required **(3)**	9	9	9	105,000
B760	114	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct basis; SAP review required; credit $78,750 toward new active ingredient application that follows	12	12	12	131,250
B761	115	Experimental Use Permit application; new active ingredient; non-food/feed or crop destruct; no SAP review required; credit $78,750 toward new active ingredient application that follows	7	7	7	78,750

B770	116	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; SAP review required; credit $105,000 toward new active ingredient application that follows	15	15	15	157,500
B771	117	Experimental Use Permit application; new active ingredient; establish temporary tolerance or tolerance exemption; no SAP review required; credit $105,000 toward new active ingredient application that follows	10	10	10	105,000
B772	118	Amend or extend Experimental Use Permit; minor changes to experimental design; established temporary tolerance or tolerance exemption is unaffected	3	3	3	10,500
B773	119	Amend or extend existing Experimental Use Permit; minor changes to experimental design; extend established temporary tolerance or tolerance exemption	5	5	5	26,250
B860	120	Amend Experimental Use Permit; first food use or major revision of experimental design	6	6	6	10,500
B780	121	New active ingredient; non-food/feed; no SAP review required **(4)**	12	12	12	131,250
B790	122	New active ingredient; Non-food/feed; SAP review required **(4)**	18	18	18	183,750
B800	123	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; no SAP review required **(4)**	12	12	12	210,000
B810	124	New active ingredient; establish permanent tolerance or tolerance exemption based on temporary tolerance or tolerance exemption; SAP review required **(4)**	18	18	18	262,500
B820	125	New active ingredient; establish tolerance or tolerance exemption; no SAP review required **(4)**	15	15	15	262,500
B840	126	New active ingredient; establish tolerance or tolerance exemption; SAP review required **(4)**	21	21	21	315,000
B830	127	New active ingredient; Experimental Use Permit application submitted simultaneously; establish tolerance or tolerance exemption; no SAP review required **(4)**	15	15	15	315,000
B850	128	New active ingredient; Experimental Use Permit requested simultaneously; establish tolerance or tolerance exemption; SAP review required **(4)**	21	21	21	367,500
B851	129	New active ingredient; different genetic event of a previously approved active ingredient; same crop; no tolerance action required; no SAP review required	9	9	9	105,000
B852	130	New active ingredient; different genetic event of a previously approved active ingredient; same crop; no tolerance action required; SAP review required	9	9	9	157,500
B870	131	New use **(3)**	9	9	9	31,500
B880	132	New product; no SAP review required **(5)**	9	9	9	26,250

B881	133	New product; SAP review required **(5)**	15	15	15	78,750
B890	134	Amendment; seed production to commercial registration; no SAP review required	9	9	9	52,500
B891	135	Amendment; seed production to commercial registration; SAP review required	15	15	15	105,000
B900	136	Amendment (except #B890); No SAP review required; (e.g., new IRM requirements that are applicant initiated; or amending a conditional registration to extend the registration expiration date with additional data submitted) **(2)**	6	6	6	10,500
B901	137	Amendment (except #B890); SAP review required **(2)**	12	12	12	63,000
B902	138	PIP Protocol review	3	3	3	5,250
B903	139	Inert ingredient tolerance exemption; e.g., a marker such as NPT II; reviewed in BPPD	6	6	6	52,500
B904	140	Import tolerance or tolerance exemption; processed commodities/food only	9	9	9	105,000

FOOTNOTES

(1)	All uses (food and/or non-food) included in any original application or petition for a new active ingredient or a first food use that otherwise satisfy the conditions for the category are covered by the base fee for that application.
(2)	EPA-initiated amendments shall not be charged fees. Fast-track amendments handled by the Antimicrobials Division are to be completed within the FIFRA stated timelines listed in Section 3(h) and are not subject to PRIA fees. Label amendments submitted by notification under PR Notices, such as PR Notice 95-2 and PR Notice 98-10, continue under PR Notice timelines and are not subject to PRIA fees.
(3)	Example: Transfer existing PIP trait by traditional breeding, such as from field corn to sweet corn.
(4)	May be either a registration for seed increase or a full commercial registration. If a seed increase registration is granted first, full commercial registration is obtained using B890.
(5)	Example: Stacking PIP traits within a crop using traditional breeding techniques.